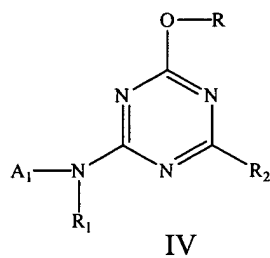
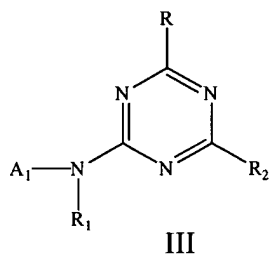
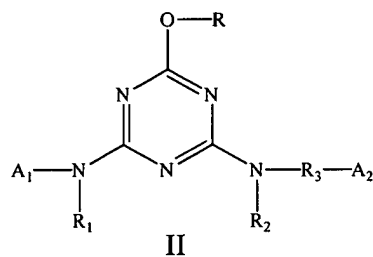
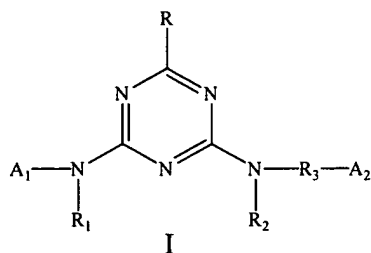


ABSTRACT

The invention is directed to compounds of Formulae I, II, III or IV:



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wherein R, R₁, R₂, R₃, A₁ and A₂ are set forth in the specification, as well as solvates, hydrates, tautomers or pharmaceutically acceptable salts thereof, that inhibit protein tyrosine kinases, especially VEGFR-2 (KDR), c-fms, c-met and tie-2 kinases. The invention is also directed toward methods of preparation of the compounds of Formulae I, II, III and IV.

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